AMENDMENTS TO THE CLAIMS

1-6. (Cancelled)

7. (Currently amended) The A percutaneous absorption preparation according to claim 17, wherein the compound having a melatonin receptor agonist activity is comprising (S)-N-[2-(1,6,7,8-tetrahydro-2H-indeno [5,4-b]furan-8-yl)ethyl]acetamide, lauric diethanolamide, and optionally one or more members selected from fatty acid esters and polyhydric alcohols.

8-19. (Cancelled)

- **20.** (Currently amended) The A percutaneous absorption preparation according to claim 17 comprising (S)-N-[2-(1,6,7,8-tetrahydro-2H-indeno[5,4-b]furan-8-yl)ethyl]acetamide, isopropyl myristate, polyethylene glycol and lauric diethanolamide.
- 21. (Currently amended) The A percutaneous absorption preparation according to claim 17 which comprising a compound having a melatonin receptor agonist activity, lauric diethanolamide and optionally one or more members selected from fatty acid esters and polyhydric alcohols, wherein the compound having a melatonin receptor agonist activity is a compound represented by the formula:

$$\begin{array}{c|c}
R^{2} \\
N \\
R^{1}
\end{array}$$

$$\begin{array}{c|c}
R^{1} \\
R^{2} \\
R^{3}
\end{array}$$

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wherein, R¹ represents an optionally substituted hydrocarbon group, an optionally substituted amino group or an optionally substituted heterocyclic group;

R² represents a hydrogen atom or an optionally substituted hydrocarbon group;

R³ represents a hydrogen atom, an optionally substituted hydrocarbon group or an optionally substituted heterocyclic group;

X represents CHR⁴, NR⁴, O or S in which R⁴ represents a hydrogen atom or an optionally substituted hydrocarbon group;

Y represents C, CH or N, provided that when X is CH₂, Y is C or CH;

represents a single bond or a double bond;

ring A represents an optionally substituted, 5- to 7-membered oxygen-containing heterocyclic ring;

ring B represents an optionally substituted benzene ring; and

m represents an integer of 1 to 4;

or a salt thereof, wherein the percutaneous absorption preparation is a skin plaster or a skin patch which is applied and/or attached to the skin.

22-32. (Cancelled)

33. (Currently amended) The A percutaneous absorption preparation according to claim 32, wherein the filler is silicon dioxide. comprising a compound having a melatonin receptor agonist activity, lauric diethanolamide and optionally one or more members selected from fatty acid esters and polyhydric alcohols, wherein the compound having a melatonin receptor agonist activity is a compound represented by the formula:

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$$\begin{array}{c|c}
R^{2} \\
N \\
N \\
R^{1}
\end{array}$$

$$\begin{array}{c|c}
R^{1} \\
CH_{2})_{m} & O \\
X
\end{array}$$

wherein, R¹ represents an optionally substituted hydrocarbon group, an optionally substituted amino group or an optionally substituted heterocyclic group;

R² represents a hydrogen atom or an optionally substituted hydrocarbon group;

R³ represents a hydrogen atom, an optionally substituted hydrocarbon group or an optionally substituted heterocyclic group;

X represents CHR⁴, NR⁴, O or S in which R⁴ represents a hydrogen atom or an optionally substituted hydrocarbon group;

Y represents C, CH or N, provided that when X is CH₂, Y is C or CH;

represents a single bond or a double bond;

ring A represents an optionally substituted, 5- to 7-membered oxygen-containing heterocyclic ring;

ring B represents an optionally substituted benzene ring; and

m represents an integer of 1 to 4;

or a salt thereof, wherein the percutaneous absorption preparation is contained in a skin contact member comprising silicon dioxide.

34-38. (Cancelled)

39. (Currently amended) A method of treating diseases related to melatonin, which comprises administrating the percutaneous absorption preparation according to claim 17 to a patient with a melatonin related disease[[.]] a percutaneous absorption preparation comprising a compound having a melatonin receptor agonist activity, lauric diethanolamide and optionally one or more members selected from fatty acid esters and polyhydric alcohols, wherein the compound having a melatonin receptor agonist activity is a compound represented by the formula:

$$\begin{array}{c|c}
R^{2} \\
N \\
N \\
R^{1}
\end{array}$$

$$\begin{array}{c|c}
R^{1} \\
R^{2} \\
N \\
R^{3}
\end{array}$$

wherein, R¹ represents an optionally substituted hydrocarbon group, an optionally substituted amino group or an optionally substituted heterocyclic group;

R² represents a hydrogen atom or an optionally substituted hydrocarbon group;

R³ represents a hydrogen atom, an optionally substituted hydrocarbon group or an optionally substituted heterocyclic group;

X represents CHR⁴, NR⁴, O or S in which R⁴ represents a hydrogen atom or an optionally substituted hydrocarbon group;

Y represents C, CH or N, provided that when X is CH₂, Y is C or CH;

represents a single bond or a double bond;

ring A represents an optionally substituted, 5- to 7-membered oxygen-containing heterocyclic ring;

ring B represents an optionally substituted benzene ring; and m represents an integer of 1 to 4;
or a salt thereof.

40. (Currently amended) A method for percutaneous absorption of a compound having a melatonin receptor agonist activity, which comprises administering the percutaneous absorption preparation according to claim 17 to a patient with a melatonin related disease[[.]] a percutaneous absorption preparation comprising a compound having a melatonin receptor agonist activity, lauric diethanolamide and optionally one or more members selected from fatty acid esters and polyhydric alcohols, wherein the compound having a melatonin receptor agonist activity is a compound represented by the formula:

$$\begin{array}{c|c}
R^{2} \\
N \\
R^{1}
\end{array}$$

$$\begin{array}{c|c}
R^{1} \\
CH_{2})_{m} & 0 \\
Y \\
X
\end{array}$$

wherein, R¹ represents an optionally substituted hydrocarbon group, an optionally substituted amino group or an optionally substituted heterocyclic group;

R² represents a hydrogen atom or an optionally substituted hydrocarbon group;

R³ represents a hydrogen atom, an optionally substituted hydrocarbon group or an optionally substituted heterocyclic group;

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X represents CHR⁴, NR⁴, O or S in which R⁴ represents a hydrogen atom or an optionally substituted hydrocarbon group;

Y represents C, CH or N, provided that when X is CH₂, Y is C or CH;

represents a single bond or a double bond;

ring A represents an optionally substituted, 5- to 7-membered oxygen-containing heterocyclic ring:

ring B represents an optionally substituted benzene ring; and m represents an integer of 1 to 4;

or a salt thereof.

41. (Cancelled)

42. (**Previously presented**) The method according to claim 39, wherein the percutaneous absorption preparation is affixed between about 6 hours before bedtime to just before bedtime.